## What is claimed is:

1. A method of preventing or treating atherosclerosis or restenosis in a mammal, comprising administering to said mammal an effective amount of a compound selected from the group consisting of structure Formula VI, Formula VII, Formula VIII and Formula IX,

# wherein Formula VI is:

or a pharmaceutically acceptable salt thereof wherein,  $\mathbf{A}^{\text{VI}}$  is

- a) Cl,
- b) Br,
- c) CN,
- d)  $NO_2$ , or
- e) F;

 $R^{VI-1}$  is

- a)  $R^{VI-5}$ , or
- b)  $SO_2R^{VI-9}$

 $R^{\text{VI-2}},\ R^{\text{VI-3}}$  and  $R^{\text{VI-4}}$  may be the same or different and are selected from the group consisting of:

- a) H,
- b) halo<sup>VI</sup>,
- c) aryl<sup>VI</sup>,
- d)  $S(0)_m R^{VI-6}$ ,
- e) (C=O)  $R^{VI-6}$ ,
- f) (C=O)  $OR^{VI-9}$ ,
- g) cyano,

- h)  $\text{het}^{\text{VI}}$ , wherein said  $\text{het}^{\text{VI}}$  is bound via a carbon atom,
- i)  $OR^{VI-10}$ ,
- j) Ohet<sup>VI</sup>,
- k)  $NR^{VI-7}R^{VI-8}$
- 1)  $SR^{VI-10}$ ,
- m) Shet<sup>VI</sup>,
- n) NHCOR<sup>VI-12</sup>,
- o)  $NHSO_2R^{VI-12}$ ,
- p)  $C_{1-7}$ alkyl which may be partially unsaturated and optionally substituted by one or more substituents of the group  $R^{VI-11}$ ,  $OR^{VI-13}$ ,  $SR^{VI-10}$ ,  $SR^{VI-13}$ ,  $NR^{VI-7}R^{VI-8}$ , halo,  $(C=0)C_{1-7}$ alkyl, or  $SO_mR^{VI-9}$ , and
- q)  $R^{VI-3}$  together with  $R^{VI-2}$  or  $R^{VI-4}$  form a carbocyclic or  $^{VI-}$ het which may be optionally substituted by  $NR^{VI-7}R^{VI-8}$ , or  $C_{1-7}$ alkyl which may be optionally substituted by  $OR^{VI-14}$ ;

## $R^{VI-5}$ is

- a)  $(CH_2CH_2O)_{i}R^{VI-10}$ ,
- b)  $C_{1-7}$ alkyl which may be partially unsaturated and is optionally substituted by one or more substituents selected from a group consisting of  $NR^{VI-7}R^{VI-8}$ ,  $R^{VI-11}$ ,  $SO_mR^{VI-9}$ , or  $OC_{2-4}$ alkyl which may be further substituted by het<sup>VI</sup>,  $OR^{VI-10}$ , or  $NR^{VI-7}R^{VI-8}$ , or
- c)  $C_{3-8}$ cycloalkyl which may be partially unsaturated and optionally substituted by one or more substituents selected from a group consisting of  $R^{VI-11}$ ,  $NR^{VI-7}R^{VI-8}$ ,  $SO_m^{VI}R^{VI-9}$ , or  $C_{1-7}$ alkyl optionally substituted by  $R^{VI-11}$ ,  $NR^{VI-7}R^{VI-8}$ , or  $SO_m^{VI}R^{VI-9}$ ;

# $R^{VI-6}$ is

- a)  $C_{1-7}$ alkyl,
- b)  $NR^{VI-7}R^{VI-8}$ ,

- c) aryl<sup>VI</sup>, or
- d) het<sup>vI</sup>, wherein said het<sup>VI</sup> is bound via a carbon atom;

 $R^{VI-7}$  and  $R^{VI-8}$  are independently

- a) H,
- b) aryl<sup>VI</sup>,
- c)  $C_{1-7}$ alkyl which may be partially unsaturated and is optionally substituted by one or more substituents selected from a group consisting of aryl<sup>VI</sup>,  $NR^{VI-10}R^{VI-10}$ ,  $R^{VI-11}$ ,  $SO_mR^{VI-9}$ ,  $CONR^{VI-10}R^{VI-10}$ , or halo, or;
- d)  $C_{3-8}$ cycloalkyl which may be partially unsaturated and optionally substituted by one or more substituents selected from a group consisting of  $R^{VI-11}$ ,  $NR^{VI-7}R^{VI-8}$ ,  $SO_m^{VI}R^{VI-9}$ , or  $C_{1-7}$ alkyl optionally substituted by  $R^{VI-11}$ ,  $NR^{VI-7}R^{VI-8}$ , or  $SO_m^{VI}R^{VI-9}$ , or
- e)  $R^{VI-7}$  and  $R^{VI-8}$  together with the nitrogen to which they are attached form a het  $^{VI}$ ;

 $R^{VI-9}$  is

- a) aryl<sup>VI</sup>,
- b) het<sup>VI</sup>,
- c)  $C_{3-8}$ cycloalkyl,
- d) methyl, or
- e)  $C_{2-7}$ alkyl which may be partially unsaturated and is optionally substituted by one or more substituents selected from a group consisting of  $NR^{VI-10}R^{VI-10}$ ,  $R^{VI-11}$ , SH,  $CONR^{VI-10}R^{VI-10}$ , or halo;

 $R^{VI-10}$  is

- a) H,
- b) methyl, or
- c) C<sub>2-7</sub>alkyl optionally substituted by OH;

 $R^{VI-11}$  is

- a)  $OR^{VI-10}$ ,
- b) Ohet<sup>VI</sup>,

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Oaryl<sup>VI</sup>,
       c)
             CO_2R^{VI--10},
       d)
             het<sup>VI</sup>,
       e)
             vi-arvl<sup>vi</sup>,
       f)
       g)
             CN, or
              C_{3-8}cycloalkyl which may be partially
       h)
              unsaturated and optionally substituted by one
              or more substituents selected from a group
              consisting of R^{VI-11}, NR^{VI-7}R^{VI-8}, SO_m^{IV}R^{VI-9}, or
              C_{1-7}alkyl optionally substituted by R^{VI-11}, NR^{VI-7}
              R^{VI-8}, or SO_mR^{VI-9};
R^{VI-12} is
       a)
              Η,
             het<sup>VI</sup>,
       b)
             aryl<sup>VI</sup>,
       C)
       d)
             C_{3-8}cycloalkyl,
             methyl, or
       e)
              C_{2-7}alkyl optionally substituted by NR^{VI-7}R^{VI-8}
       f)
              or R<sup>VI-11</sup>;
R^{VI-13} is
              (P=0) (OR^{VI-14})_{2}
       a)
              CO(CH_2)_n^{IV}CON(CH_3) - (CH_2)_nSO_3^{IV}M^{VI+}
       b)
              an amino<sup>VI</sup> acid,
       C)
              C(=0) aryl<sup>VI</sup>,
       d)
              C(=0)C_{1-7}alkyl optionally substituted by NR^{VI-7}
       e)
              R^{VI-8}, aryl<sup>VI</sup>, het<sup>VI</sup>, CO_2H, or O(CH_2)_nCO_2R^{VI-14}, or
              C (=0) NR^{VI-7} R^{VI-8}
       f)
R^{VI-14} is
       a)
            H, or
            C_{1-7}alkyl;
       b)
each i<sup>VI</sup> is independently 2, 3, or 4;
each n^{VI} is independently 1, 2, 3, 4 or 5;
each m<sup>VI</sup> is independently 0, 1, or 2;
M^{VI} is sodium, potassium, or lithium;
aryl is a phenyl radical or an ortho-fused bicyclic
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carbocyclic radical wherein at least one ring is aromatic;

wherein any aryl<sup>VI</sup> is optionally substituted with one or more substituents selected from the group consisting of halo, OH, cyano,  $CO_2R^{VI-14}$ ,  $CF_3$ ,  $C_{1-6}$ alkoxy, and  $C_{1-6}$  alkyl which maybe further substituted by one to three  $SR^{VI-14}$ ,  $NR^{VI-14}R^{VI-14}$ ,  $OR^{VI-14}$ , or  $CO_2R^{VI-14}$ ;

het<sup>vI</sup> is a four- (4), five- (5), six- (6), or seven- (7) membered saturated or unsaturated heterocyclic ring having 1, 2, or 3 heteroatoms selected from the group consisting of oxygen, sulfur, and nitrogen, which is optionally fused to a benzene ring, or any bicyclic heterocycle group;

wherein any het<sup>VI</sup> is optionally substituted with one or more substituents selected from the group consisting of halo, OH, cyano, phenyl,  $CO_2R^{VI-14}$ ,  $CF_3$ ,  $C_{1-6}$ alkoxy, oxo, oxime, and  $C_{1-6}$  alkyl which maybe further substituted by one to three  $SR^{VI-14}$ ,  $NR^{VI-14}R^{VI-14}$ ,  $OR^{VI-14}$ , or  $CO_2R^{VI-14}$ ;

wherein Formula VII is

VII

or a pharmaceutically acceptable salt thereof, wherein  $% \left( 1,...,n\right) =\left( 1,...,n\right)$ 

A<sup>VII</sup> is

- a) Cl,
- b) Br,
- c) CN,
- d)  $NO_2$ , or
- e) F;

## $R^{VII-1}$ is

- a) aryl<sup>VII</sup>,
- b)  $S(0)_{m}^{VII}R^{VII-6}$
- c) (C=O) $R^{VII-6}$ , with the proviso that if  $R^{VII-6}$  is  $NR^{VII-7}R^{VII-8}$ , then  $R^{VII-7}$  and  $R^{VII-8}$  do not both equal H,
- d) (C=O)  $OR^{VII-9}$ ,
- e) cyano,
- f)  $\text{het}^{\text{VII}}$ , wherein said  $\text{het}^{\text{VII}}$  is bound via a carbon atom,
- g) Ohet VII,
- h)  $NR^{VII-7}R^{VII-8}$  with the proviso that  $R^{VII-7}$  and  $R^{VII-8}$  do not both equal H,
- i) SR<sup>VII-10</sup>,
- j) Shet<sup>VII</sup>,
- k) NHCOR<sup>VII-12</sup>,
- 1)  $NHSO_2R^{VII-12}$ ,
- m)  $C_{1-7}$ alkyl which is partially unsaturated and optionally substituted by one or more substituents of the group  $R^{VII-11}$ ,  $OR^{VII-13}$ ,  $SR^{VII-10}$ ,  $SR^{VII-13}$ ,  $NR^{VII-7}R^{VII-8}$ , halo,  $(C=O)C_{1-7}$ alkyl, or  $SO_mR^{VII-9}$ , or
- n)  $C_{1-7}$ alkyl which is substituted by one or more substituents of the group  $R^{VII-11}$ ,  $OR^{VII-13}$ ,  $SR^{VII-10}$ ,  $SR^{VII-13}$ ,  $NR^{VII-7}R^{VII-8}$ , halo, (C=O) $C_{1-7}$ alkyl, or  $SO_m^{VII}R^{VII-9}$ ;

## $R^{VII-2}$ is

- a) H,
- b) halo,
- c) aryl<sup>VII</sup>,
- d)  $S(0)_m^{VII}R^{VII-6}$ ,
- e)  $(C=0) R^{VII-6}$ ,
- f) (C=O)  $OR^{VII-9}$ ,
- g) cyano,

- h) het<sup>VII</sup>, wherein said het<sup>VII</sup> is bound via a carbon atom,
- i) OR $^{VII-10}$ ,
- j) Ohet<sup>VII</sup>,
- k)  $NR^{VII-7}R^{VII-8}$
- 1)  $SR^{VII-10}$ ,
- m) Shet<sup>VII</sup>,
- n) NHCOR<sup>VII-12</sup>,
- o) NHSO<sub>2</sub>R<sup>VII-12</sup>, or
- p)  $C_{1-7}$ alkyl which may be partially unsaturated and optionally substituted by one or more substituents of the group  $R^{VII-11}$ ,  $OR^{VII-13}$ ,  $SR^{VII-10}$ ,  $SR^{VII-13}$ ,  $NR^{VII-7}R^{VII-8}$ , halo,  $(C=0)C_{1-7}$ alkyl, or  $SO_m^{VII}R^{VII-9}$ , or
- q)  $R^{VII-1}$  together with  $R^{VII-2}$  form a carbocyclic or het<sup>VII</sup> which may be optionally substituted by  $NR^{VII-7}R^{VII-8}$ , or  $C_{1-7}$ alkyl which may be optionally substituted by  $OR^{VII-14}$ ;

## $R^{VII-6}$ is

- a)  $C_{1-7}$ alkyl,
- b) NR<sup>VII-7</sup>R<sup>VII-8</sup>
- c) aryl<sup>VII</sup>, or
- d) het $^{VII}$ , wherein said het $^{VII}$  is bound via a carbon atom;

## $R^{VII-7}$ and $R^{VII-8}$ are independently

- a) H,
- b) aryl<sup>VII</sup>,
- c)  $C_{1-7}$ alkyl which may be partially unsaturated and is optionally substituted by one or more substituents selected from  $NR^{VII-10}R^{VII-10}$ ,  $R^{VII-11}$ ,  $SO_mR^{VII-9}$ ,  $CONR^{VII-10}R^{VII-10}$ , or halo, or,
- d)  $R^{VII-7}$  and  $R^{VII-8}$  together with the nitrogen to which they are attached form a het<sup>VII</sup>;

#### R<sup>VII-9</sup> is

a) aryl<sup>VII</sup>,

- b) het<sup>VII</sup>,
- c)  $C_{3-8}$ cycloalkyl,
- d) methyl, or
- e)  $C_{2-7}$ alkyl which may be partially unsaturated and is optionally substituted by one or more substituents selected from  $NR^{VII-10}R^{VII-10}$ ,  $R^{VII-11}$ , SH,  $CONR^{VII-10}R^{VII-10}$ , or halo;

#### $R^{VII-10}$ is

- a) H,
- b) methyl, or
- c)  $C_{2-7}$ alkyl optionally substituted by OH;

#### $R^{VII-11}$ is

- a)  $OR^{VII-10}$ ,
- b) Ohet VII,
- c) Oaryl<sup>VII</sup>,
- d)  $CO_2R^{VII-10}$
- e) het<sup>VII</sup>,
- f) aryl<sup>VII</sup>,
- g) CN, or
- h)  $C_{3-8}$ cycloalkyl which may be partially unsaturated and optionally substituted by one or more substituents seleted from a group consisting of  $R^{VII-11}$ ,  $NR^{VII-7}R^{VII-8}$ ,  $SO_m^{VII}R^{VII-9}$ , or  $C_{1-7}$ alkyl optionally substituted by  $R^{VII-11}$ ,  $NR^{VII-7}R^{VII-8}$ , or  $SO_mR^{VII-9}$ ;

#### $R^{VII-12}$ is

- a) H,
- b) het<sup>VII</sup>,
- c) aryl<sup>VII</sup>,
- d) C<sub>3-8</sub>cycloalkyl,
- e) methyl, or
- f)  $C_{2-7}$ alkyl optionally substituted by  $NR^{VII-7}R^{VII-8}$  or  $R^{VII-11}$ ;

## $R^{VII-13}$ is

a)  $(P=0) (OR^{VII-14})_2$ ,

- b)  $CO(CH_2)_nCON(CH_3) (CH_2)_nSO_3^-M^+$ ,
- c) an amino acid,
- d) C(=0) aryl<sup>VII</sup>, or
- e)  $C(=0)C_{1-7}alkyl$  optionally substituted by  $NR^{VII-7}R^{VII-8}$ ,  $aryl^{VII}$ ,  $het^{VII}$ ,  $CO_2H$ , or  $O(CH_2)_n^{VII}CO_2R^{VII-14}$ ;

#### $R^{VII-14}$ is

- a) H, or
- b)  $C_{1-7}$ alkyl;

each  $n^{VII}$  is independently 1, 2, 3, 4 or 5; each  $m^{VII}$  is independently 0, 1, or 2;  $M^{VII}$  is sodium, potassium, or lithium;

- aryl<sup>VII</sup> is a phenyl radical or an ortho-fused bicyclic carbocyclic radical wherein at least one ring is aromatic;
- wherein any aryl<sup>VII</sup> is optionally substituted with one or more substituents selected from the group consisting of halo, OH, cyano,  $CO_2R^{VII-14}$ ,  $CF_3$ ,  $C_{1-6}$ alkoxy, and  $C_{1-6}$  alkyl which may be further substituted by one to three  $SR^{VII-14}$ ,  $NR^{VII-14}R^{VII-14}$ ,  $OR^{VII-14}$ , or  $CO_2R^{VII-14}$  groups;
- het<sup>VII</sup> is a four- (4), five- (5), six- (6), or seven- (7) membered saturated or unsaturated heterocyclic ring having 1, 2, or 3 heteroatoms selected from the group consisting of oxygen, sulfur, and nitrogen, which is optionally fused to a benzene ring, or any bicyclic heterocycle group;
- wherein any het<sup>VII</sup> is optionally substituted with one or more substituents selected from the group consisting of halo, OH, cyano, phenyl,  $CO_2R^{VII-14}$ ,  $CF_3$ ,  $C_{1-6}$ alkoxy, oxo, oxime, and  $C_{1-6}$  alkyl which may be further substituted by one to three  $SR^{VII-14}$ ,  $NR^{VII-14}R^{VII-14}$ ,  $OR^{VII-14}$ , or  $CO_2R^{VII-14}$  groups;

wherein Formula VIII is

and pharmaceutically acceptable salts thereof, wherein

## $A^{VIII}$ is

- a) Cl,
- b) Br,
- c) CN,
- d)  $NO_2$ , or
- e) F;

#### $R^{VIII-1}$ is

- a)  $R^{VIII-5}$ ,
- b)  $NR^{VIII-7}R^{VIII-8}$ , or
- c)  $SO_2R^{VIII-9}$ ;

#### $R^{VIII-2}$ is

- a) aryl<sup>VIII</sup>,
- b) het<sup>VIII</sup>,
- c) SOmR<sup>VIII-6</sup>,
- d)  $OC_{2-7}$  alkyl substituted by OH,
- e)  $SC_{2-7}$  alkyl substituted by OH, or
- f)  $C_{2-8}$  alkyl which is partially unsaturated and is optionally substituted by one or more substituents selected from  $R^{VIII-11}$ ,  $OR^{VIII-13}$ ,  $SR^{VIII-13}$ ,  $NR^{VIII-7}R^{VIII-8}$ , halo,  $(C=0)C_{1-7}$  alkyl or  $SO_m^{VIII}R^{VIII-9}$ ;

with the proviso that when  $R^{VIII-1}=R^{VIII-5}=$  (CH<sub>2</sub>CH<sub>2</sub>O)<sub>i</sub> $^{VIII}R^{VIII-10}$ , then  $R^{VIII-2}$  may additionally represent

- a) H,
- b) halo,
- $(C=0) R^{VIII-6}$

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d)
                (C=O)OR^{VIII-9},
        e)
               cyano,
               OR<sup>VIII-10</sup>,
        f)
               het<sup>VIII</sup>,
        g)
               NR<sup>VIII-7</sup>R<sup>VIII-8</sup>,
        h)
               SR<sup>VIII-10</sup>,
        i)
               het<sup>VIII</sup>,
        j)
               NHCOR VIII-12,
        k)
               NHSO_2R^{VIII-12}, or
        1)
               R^{VIII-2} together with R^{VIII-3} or R^{VIII-4} form a
        m)
               carbocyclic or het VIII which may be optionally
               substituted by NR^{VIII-7}R^{VIII-8}, or C_{1-7}alkyl which
               may be optionally substituted by OR^{VIII-14};
R^{\text{VIII-3}} and R^{\text{VIII-4}} are independently:
        a)
               Η,
       b)
               halo,
               aryl^{VIII},
        C)
               S(0)_{m}^{viii}R^{viii-6},
        d)
               (C=O) R^{VIII-6},
        e)
               (C=O) OR<sup>VIII-9</sup>,
        f)
        g)
               cyano,
               het viii, wherein said het viii is bound via a
        h)
                carbon atom,
               OR<sup>VIII-10</sup>,
        i)
               Ohet VIII,
        j)
               NRVIII-7RVIII-8,
        k)
               SR<sup>VIII-10</sup>,
        1)
               Shet<sup>VIII</sup>,
        m)
               NHCOR VIII-12,
        n)
               NHSO<sub>2</sub>R<sup>VIII-12</sup>,
        0)
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optionally substituted by one or more

 $C_{1-7}$ alkyl which may be partially unsaturated and

p)

substituents of the group  $R^{VIII-11}$ ,  $OR^{VIII-13}$ ,  $SR^{VIII-10}$ ,  $SR^{VIII-13}$ ,  $NR^{VIII-7}R^{VIII-8}$ , halo, (C=O)C<sub>1-7</sub>alkyl, or  $SO_m^{VIII}RVIII^{-9}$ , or

q)  $R^{VIII-4}$  together with  $R^{VIII-3}$  form a carbocyclic or het which may be optionally substituted by  $NR^{VIII-7}R^{VIII-8}$ , or  $C_{1-7}$ alkyl which may be optionally substituted by  $OR^{VIII-14}$ ;

#### $R^{VIII-5}$ is

- a)  $(CH_2CH_2O)_iR^{VIII-10}$ ,
- b) het<sup>VIII</sup>, wherein said het<sup>VIII</sup> is bound via a carbon atom,
- c) aryl<sup>VIII</sup>,
- d)  $C_{1-7}$ alkyl which may be partially unsaturated and is optionally substituted by one or more substituents selected from  $NR^{VIII-7}R^{VIII-8}$ ,  $R^{VIII-11}$ ,  $SO_mR^{VIII-9}$ , or  $OC_{2-4}$ alkyl which may be further substituted by het  $V^{VIII}$ ,  $OR^{VIII-10}$ , or  $V^{VIII-7}R^{VIII-8}$ , or
- e)  $C_{3-8}$ cycloalkyl which may be partially unsaturated and optionally substituted by one or more substituents selected from  $R^{VIII-11}$ ,  $NR^{VIII-7}R^{VIII-8}$ ,  $SO_m^{VIII}R^{VIII-9}$ , or  $C_{1-7}$ alkyl optionally substituted by  $R^{VIII-11}$ ,  $NR^{VIII-7}R^{VIII-8}$ , or  $SO_m^{VIII}R^{VIII-9}$ ;

#### R<sup>VIII-6</sup> is

- a)  $C_{1-7}$ alkyl,
- b)  $NR^{VIII-7}R^{VIII-8}$
- c) aryl<sup>VIII</sup>, or
- d) het<sup>VIII</sup>, wherein said het<sup>VIII</sup> is bound via a carbon atom;

 $R^{\text{VIII-7}}$  and  $R^{\text{VIII-8}}$  are independently

a) H,

- b) aryl<sup>VIII</sup>,
- c)  $C_{1-7}$ alkyl which may be partially unsaturated and is optionally substituted by one or more substituents selected from  $NR^{VIII-10}R^{VIII-10}$ ,  $R^{VIII-11}$ ,  $SO_m^{VIII}R^{VIII-9}$ ,  $CONR^{VIII-10}R^{VIII-10}$ , or halo, or,
- d) R<sup>VIII-7</sup> and R<sup>VIII-8</sup> together with the nitrogen to which they are attached form a het<sup>VIII</sup>;

# R<sup>VIII-9</sup> is

- a) aryl<sup>VIII</sup>,
- b) het<sup>VIII</sup>,
- c)  $C_{3-8}$ cycloalkyl,
- d) methyl, or
- e)  $C_{2-7}$ alkyl which may be partially unsaturated and is optionally substituted by one or more substituents selected from  $NR^{VIII-10}R^{VIII-10}$ ,  $R^{VIII-11}$ , SH,  $CONR^{VIII-10}R^{VIII-10}$ , or halo;

## R<sup>VIII-10</sup> is

- a) H,
- b) methyl, or
- c)  $C_{2-7}$ alkyl optionally substituted by OH;

#### $R^{VIII-11}$ is

- a) OR<sup>VIII-10</sup>,
- b) Ohet VIII,
- c) Oaryl<sup>VIII</sup>,
- d)  $CO_2R^{VIII-10}$ ,
- e) het<sup>VIII</sup>,
- f) aryl<sup>VIII</sup>, or
- g) CN;

# $R^{VIII-12}$ is

a) H,

- b) het<sup>VIII</sup>,
- c) aryl<sup>VIII</sup>,
- d)  $C_{3-8}$ cycloalkyl,
- e) methyl, or
- f)  $C_{2-7}$ alkyl optionally substituted by  $NR^{VIII-7}R^{VIII-8}$  or  $R^{VIII-11}$ :

## $R^{VIII-13}$ is

- a)  $(P=0) (OR^{14})_{2}$
- b)  $CO(CH_2)_n^{VIII}CON(CH_3) (CH_2)_n^{VIII}SO_3^{-M^+}$ ,
- c) an amino acid,
- d) C(=0) aryl<sup>VIII</sup>, or
- e)  $C(=0) C_{1-7}alkyl$  optionally substituted by  $NR^{VIII-7}R^{VIII-8}$ ,  $aryl^{VIII}$ ,  $het^{VIII}$ ,  $CO_2H$ , or  $O(CH_2)_n^{VIII}CO_2R^{VIII-14}$ ;

### R<sup>VIII-14</sup> is

- a) H, or
- b)  $C_{1-7}$ alkyl;

each i<sup>VIII</sup> is independently 2, 3, or 4;

each n<sup>VIII</sup> is independently 1, 2, 3, 4 or 5;

each m<sup>VIII</sup> is independently 0, 1, or 2;

M<sup>VIII</sup> is sodium, potassium, or lithium;

- aryl viii is a phenyl radical or an ortho-fused bicyclic carbocyclic radical wherein at least one ring is aromatic;
- wherein any aryl<sup>VIII</sup> is optionally substituted with one or more substituents selected from halo, OH, cyano,  $CO_2R^{VIII-14}$ ,  $CF_3$ ,  $C_{1-6}$ alkoxy, and  $C_{1-6}$  alkyl which may be further substituted by one to three  $SR^{VIII-14}$ ,  $NR^{VIII-14}R^{VIII-14}$ ,  $OR^{VIII-14}$ , or  $CO_2R^{VIII-14}$  groups;
- het<sup>VIII</sup> is a four- (4), five- (5), six- (6), or seven- (7) membered saturated or unsaturated heterocyclic ring having 1, 2, or 3 heteroatoms selected from the

group consisting of oxygen, sulfur, and nitrogen, which is optionally fused to a benzene ring, or any bicyclic heterocycle group;

wherein any het<sup>VIII</sup> is optionally substituted with one or more substituents selected from the group consisting of halo, OH, cyano, phenyl,  $CO_2R^{VIII-14}$ ,  $CF_3$ ,  $C_{1-6}$ alkoxy, oxo, oxime, and  $C_{1-6}$  alkyl which may be further substituted by one to three  $SR^{VIII-14}$ ,  $NR^{VIII-14}R^{VIII-14}$ ,  $OR^{VIII-14}$ , or  $CO_2R^{VIII-14}$  groups;

wherein Formula IX is

ΙX

and pharmaceutically acceptable salts thereof, wherein,

 $R^{IX-1}$  is

- a) Cl,
- b) Br,
- c) CN,
- d)  $NO_2$ , or
- e) F;

 $R^{\text{IX-2}}$ ,  $R^{\text{IX-3}}$  and  $R^{\text{IX-4}}$  are independently selected from:

- a) H,
- b) halo,
- c) aryl<sup>IX</sup>,
- d)  $S(0)_{m}^{IX}R^{IX-6}$ ,
- e) (C=O)  $R^{IX-6}$ ,
- f) (C=O)  $OR^{IX-9}$ ,
- g) cyano,
- h)  $\text{het}^{\text{IX}}$ , wherein said IX-het is bound via a carbon atom,

- i)  $OR^{IX-10}$ ,
- j) Ohet<sup>IX</sup>,
- k)  $NR^{IX-7}R^{IX-8}$
- 1)  $SR^{IX-10}$ ,
- m) Shet<sup>IX</sup>,
- n) NHCOR<sup>IX-12</sup>,
- o)  $NHSO_2R^{IX-12}$ , or
- p)  $C_{1-7}$ alkyl which may be partially unsaturated and optionally substituted by one or more substituents of the group  $R^{IX-11}$ ,  $OR^{IX-13}$ ,  $SR^{IX-10}$ ,  $SR^{IX-13}$ ,  $NR^{IX-7}R^{IX-8}$ , halo,  $(C=0)C_{1-7}$ alkyl, or  $SO_m^{IX}R^{IX-9}$ ;

# $R^{IX-6}$ is

- a)  $C_{1-7}$ alkyl,
- b)  $NR^{IX-7}R^{IX-8}$ ,
- c) aryl<sup>IX</sup>, or
- d) het<sup>IX</sup>, wherein said het<sup>IX</sup> is bound via a carbon atom;

## $R^{IX-7}$ and $R^{IX-8}$ are independently

- a) H,
- b) aryl<sup>IX</sup>,
- c)  $C_{1-7}$ alkyl which may be partially unsaturated and is optionally substituted by one or more substituents selected from  $NR^{IX-10}R^{IX-10}$ ,  $R^{IX-11}$ ,  $SO_mR^{IX-9}$ ,  $CONR^{IX-10}R^{IX-10}$ , or halo, or,
- d)  $R^{IX-7}$  and  $R^{IX-8}$  together with the nitrogen to which they are attached form a IX-het;

#### RIX-9 is

- a) aryl<sup>IX</sup>,
- b) het<sup>IX</sup>,
- c)  $C_{3-8}$ cycloalkyl,
- d) methyl, or

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e)
               C_{2-7}alkyl which may be partially unsaturated and
                is optionally substituted by one or more
                substituents selected from NR^{IX-10}R^{IX-10}, R^{IX-11},
                SH, CONR<sup>IX-10</sup>R<sup>IX-10</sup>, or halo;
R^{IX-10} is
        a)
               Η,
               methyl, or
        b)
               C_{2-7}alkyl optionally substituted by OH;
R^{IX-11} is
               OR<sup>IX-10</sup>,
        a)
       b)
               Ohet IX,
               Oaryl<sup>IX</sup>,
        C)
               CO_2R^{IX-10},
        d)
               het<sup>IX</sup>,
        e)
               aryl<sup>IX</sup>, or
        f)
               CN;
R^{IX-12} is
        a)
               Η,
               het<sup>IX</sup>,
        b)
               aryl<sup>IX</sup>,
        C)
        d)
               C_{3-8}cycloalkyl,
               methyl, or
        e)
               \text{C}_{\text{2-7}}\text{alkyl} optionally substituted by \text{NR}^{\text{IX-7}}\text{R}^{\text{IX-8}} or
        f)
                R<sup>IX-11</sup>;
R^{IX-13} is
                (P=0) (OR^{IX-14})_{2}
        a)
               CO(CH_2)_n^{IX}CON(CH_3) - (CH_2)_n^{IX}SO_3^{-M}^{IX+},
        b)
        C)
                an amino acid,
               C(=0)aryl<sup>IX</sup>, or
        d)
                C(=0)C_{1-7}alkyl optionally substituted by
               NR^{IX-7}R^{IX-8}, aryl<sup>IX</sup>, het<sup>IX</sup>, CO_2H, or O(CH_2)_nCO_2R^{IX-14};
R^{IX-14} is
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- a) H, or
- b)  $C_{1-7}$ alkyl;

each  $n^{\text{IX}}$  is independently 1, 2, 3, 4 or 5; each  $m^{\text{IX}}$  is independently 0, 1, or 2;

M<sup>IX</sup> is sodium, potassium, or lithium;

- aryl is a phenyl radical or an ortho-fused bicyclic carbocyclic radical wherein at least one ring is aromatic;
- wherein any  $\operatorname{aryl}^{\operatorname{IX}}$  is optionally substituted with one or more substituents selected from the group consisting of halo, OH, cyano,  $\operatorname{CO}_2\operatorname{R}^{\operatorname{IX}-14}$ ,  $\operatorname{CF}_3$ ,  $\operatorname{C}_{1-6}\operatorname{alkoxy}$ , and  $\operatorname{C}_{1-6}$  alkyl which may be further substituted by one to three  $\operatorname{SR}^{\operatorname{IX}-14}$ ,  $\operatorname{NR}^{\operatorname{IX}-14}\operatorname{R}^{\operatorname{IX}-14}$ ,  $\operatorname{OR}^{\operatorname{IX}-14}$ , or  $\operatorname{CO}_2\operatorname{R}^{\operatorname{IX}-14}$  groups;
- het<sup>IX</sup> is a four- (4), five- (5), six- (6), or seven- (7) membered saturated or unsaturated heterocyclic ring having 1, 2, or 3 heteroatoms selected from the group consisting of oxygen, sulfur, and nitrogen, which is optionally fused to a benzene ring, or any bicyclic heterocycle group;
- wherein any het<sup>IX</sup> is optionally substituted with one or more substituents selected from the group consisting of halo, OH, cyano, phenyl,  $CO_2R^{IX-14}$ ,  $CF_3$ ,  $C_{1-6}$ alkoxy, oxo, oxime, and  $C_{1-6}$  alkyl which may be further substituted by one to three  $SR^{IX-14}$ ,  $NR^{IX-14}R^{IX-14}$ ,  $OR^{IX-14}$ , or  $CO_2R^{IX-14}$  groups.

2. The method of claim 1, wherein the compound administered has the Formula

or a pharmaceutically acceptable salt thereof, wherein,

 $A^{\text{VI}}$  is

- a) Cl,
- b) Br,
- c) CN,
- d)  $NO_2$ , or
- e) F;

 $R^{VI-1}$  is

- a)  $R^{VI-5}$ , or
- b)  $SO_2R^{VI-9}$

 $R^{VI-2}$ ,  $R^{VI-3}$  and  $R^{VI-4}$  may be the same or different and are selected from the group consisting of:

- a) H,
- b) halo,
- c) aryl<sup>vi</sup>,
- d)  $S(0)_{m}^{VI}R^{VI-6}$ ,
- e) (C=O)  $R^{VI-6}$ ,
- f) (C=O)  $OR^{VI-9}$
- g) cyano,
- h) het  $^{\text{VI}}$ , wherein said het  $^{\text{VI}}$  is bound via a carbon atom,
- i) OR $^{VI-10}$ ,
- j) Ohet<sup>VI</sup>,
- k)  $NR^{VI-7}R^{VI-8}$
- 1)  $SR^{VI-10}$ ,

- m) Shet<sup>VI</sup>,
- n) NHCOR<sup>VI-12</sup>,
- o)  $NHSO_2R^{VI-12}$ ,
- p)  $C_{1-7}$ alkyl which may be partially unsaturated and optionally substituted by one or more substituents of the group  $R^{VI-11}$ ,  $OR^{VI-13}$ ,  $SR^{VI-10}$ ,  $SR^{VI-13}$ ,  $NR^{VI-7}R^{VI-8}$ , halo,  $(C=0)C_{1-7}$ alkyl, or  $SO_m^{VI}RVI^{-9}$ , and
- q)  $R^{VI-3}$  together with  $R^{VI-2}$  or  $R^{VI-4}$  form a carbocyclic or het which may be optionally substituted by  $NR^{VI-7}R^{VI-8}$ , or  $C_{1-7}$ alkyl which may be optionally substituted by  $OR^{VI-14}$ ;

## $R^{VI-5}$ is

- a)  $(CH_2CH_2O)_{i}^{VI}R^{VI-10}$ ,
- b)  $C_{1-7}$ alkyl which may be partially unsaturated and is optionally substituted by one or more substituents selected from a group consisting of  $NR^{VI-7}R^{VI-8}$ ,  $R^{VI-11}$ ,  $SO_m^{VI}R^{VI-9}$ , or  $OC_{2-4}$ alkyl which may be further substituted by het<sup>VI</sup>,  $OR^{VI-10}$ , or  $NR^{VI-7}R^{VI-8}$ , or
- C)  $C_{3-8}$ cycloalkyl which may be partially unsaturated and optionally substituted by one or more substituents selected from a group consisting of  $R^{VI-11}$ ,  $NR^{VI-7}R^{VI-8}$ ,  $SO_m^{VI}R^{VI-9}$ , or  $C_{1-7}$ alkyl optionally substituted by  $R^{VI-11}$ ,  $NR^{VI-7}R^{VI-8}$ , or  $SO_m^{VI}R^9$ ;

# $R^{VI-6}$ is

- a)  $C_{1-7}$ alkyl,
- b)  $NR^{VI-7}R^{VI-8}$ ,
- c) aryl<sup>VI</sup>, or
- d)  $\text{het}^{VI}$ , wherein said  $\text{het}^{VI}$  is bound via a carbon atom;

#### $R^{VI-7}$ and $R^{VI-8}$ are independently

- a) H,
- b) aryl<sup>VI</sup>,

- c)  $C_{1-7}$ alkyl which may be partially unsaturated and is optionally substituted by one or more substituents selected from a group consisting of aryl<sup>VI</sup>,  $NR^{VI-10}R^{VI-10}$ ,  $R^{VI-11}$ ,  $SO_m^{VI}R^{VI-9}$ ,  $CONR^{VI-10}R^{VI-10}$ , or halo, or;
- d)  $C_{3-8}$ cycloalkyl which may be partially unsaturated and optionally substituted by one or more substituents selected from a group consisting of  $R^{VI-11}$ ,  $NR^{VI-7}R^{VI-8}$ ,  $SO_m^{VI}R^{VI-9}$ , or  $C_{1-7}$ alkyl optionally substituted by  $R^{VI-11}$ ,  $NR^{VI-7}R^{VI-8}$ , or  $SO_m^{VI}R^{VI-9}$ , or
- e)  $R^{VI-7}$  and  $R^{VI-8}$  together with the nitrogen to which they are attached form a het<sup>VI</sup>;

# $R^{VI-9}$ is

- a) aryl<sup>VI</sup>,
- b) het<sup>VI</sup>,
- c)  $C_{3-8}$ cycloalkyl,
- d) methyl, or
- e)  $C_{2-7}$ alkyl which may be partially unsaturated and is optionally substituted by one or more substituents selected from a group consisting of  $NR^{VI-10}R^{VI-10}$ ,  $R^{VI-11}$ , SH,  $CONR^{VI-10}R^{VI-10}$ , or halo;

# $R^{VI-10}$ is

- a) H,
- b) methyl, or
- c) C<sub>2-7</sub>alkyl optionally substituted by OH;

# $R^{VI-11}$ is

- a)  $OR^{10}$ ,
- b) Ohet<sup>VI</sup>,
- c) Oaryl<sup>VI</sup>,
- d)  $CO_2R^{10}$ ,
- e) het<sup>VI</sup>,
- f) aryl<sup>VI</sup>,
- g) CN, or

h) C<sub>3-8</sub>cycloalkyl which may be partially unsaturated and optionally substituted by one or more substituents selected from a group consisting of  $R^{VI-11}$ ,  $NR^{VI-7}R^{VI-8}$ ,  $SO_m^{VI}R^{VI-9}$ , or  $C_{1-7}$ alkyl optionally substituted by  $R^{VI-11}$ ,  $NR^{VI-7}R^{VI-8}$ , or  $SO_m^{VI}R^{VI-9}$ ;  $R^{VI-12}$  is Η, a) het<sup>VI</sup>, b) C) aryl<sup>VI</sup>, d)  $C_{3-8}$ cycloalkyl, methyl, or e) C<sub>2-7</sub>alkyl optionally substituted by NR<sup>VI-7</sup>R<sup>VI-8</sup> f) or R<sup>VI-11</sup>;  $R^{VI-13}$  is  $(P=0) (OR^{VI-14})_{2}$ a)  $CO(CH_2)_n^{VI}CON(CH_3) - (CH_2)_nSO_3^{-M^{VI+}}$ , b) C) an amino acid, C(=0) ary  $l^{VI}$ , d)  $C(=0)C_{1-7}alkyl$  optionally substituted by e) NR<sup>VI-7</sup> R<sup>VI-8</sup>, aryl<sup>VI</sup>, het<sup>VI</sup>, CO<sub>2</sub>H, or  $O(CH_2)_n^{VI}CO_2R^{VI-14}$ , or  $C (=0) NR^{VI-7} R^{VI-8}$ f) R<sup>VI-14</sup> is a) H, or  $C_{1-7}$ alkyl; each i<sup>VI</sup> is independently 2, 3, or 4; each n<sup>VI</sup> is independently 1, 2, 3, 4 or 5; each m<sup>VI</sup> is independently 0, 1, or 2; M<sup>VI</sup> is sodium, potassium, or lithium; aryl VI is a phenyl radical or an ortho-fused bicyclic carbocyclic radical wherein at least one ring is aromatic;

wherein any  $\operatorname{aryl}^{\operatorname{vi}}$  is optionally substituted with one or

more substituents selected from the group consisting of halo, OH, cyano,  $CO_2R^{VI-14}$ ,  $CF_3$ ,  $C_{1-6}$ alkoxy, and  $C_{1-6}$  alkyl which maybe further substituted by one to three  $SR^{VI-14}$ ,  $NR^{VI-14}R^{VI-14}$ ,  $OR^{VI-14}$ , or  $CO_2R^{VI-14}$ ;

- het<sup>VI</sup> is a four- (4), five- (5), six- (6), or seven- (7) membered saturated or unsaturated heterocyclic ring having 1, 2, or 3 heteroatoms selected from the group consisting of oxygen, sulfur, and nitrogen, which is optionally fused to a benzene ring, or any bicyclic heterocycle group;
- wherein any het<sup>VI</sup> is optionally substituted with one or more substituents selected from the group consisting of halo, OH, cyano, phenyl,  $CO_2R^{VI-14}$ ,  $CF_3$ ,  $C_{1-6}$ alkoxy, oxo, oxime, and  $C_{1-6}$  alkyl which maybe further substituted by one to three  $SR^{VI-14}$ ,  $NR^{VI-14}R^{VI-14}$ ,  $OR^{VI-14}$ , or  $CO_2R^{VI-14}$ .
  - 3. The method of Claim 2, wherein  $A^{VI}$  is Cl.
- 4. The method of Claim 2, wherein the compound administered is selected from the group consisting of

N-(4-chlorobenzyl)-6-iodo-1-methyl-4-oxo-1,4-dihydro-3-cinnolinecarboxamide; N-(4-chlorobenzyl)-6-(3-hydroxy-1-propynyl)-1-methyl-4-oxo-1,4-dihydro-3-cinnolinecarboxamide;

N-(4-chlorobenzyl)-6-(hydroxymethyl)-1-methyl-4-oxo-1,4-dihydro-3-cinnolinecarboxamide;

N-(4-chlorobenzyl)-6-(4-hydroxy-1-butynyl)-1-methyl-4-oxo-1,4-dihydro-3-cinnolinecarboxamide;

N-(4-chlorobenzyl)-1-methyl-6-(4-morpholinylmethyl)-4- oxo-1,4-dihydro-3-cinnolinecarboxamide; N-(4-

```
chlorobenzyl) -8-\{[(1R,2R)-1-hydroxy-2-
methylcyclohexyl]ethynyl}-1-methyl-4-oxo-6-(tetrahydro-
2H-pyran-4-ylmethyl)-1,4-dihydro-3-cinnolinecarboxamide;
N-(4-chlorobenzyl)-8-(cyclopropylethynyl)-1-methyl-6-(4-
morpholinylmethyl) -4-oxo-1, 4-dihydro-3-
cinnolinecarboxamide:
N-(4-chlorobenzyl)-8-[3-(dimethylamino)-1-propynyl]-1-
methyl-6-(4-morpholinylmethyl)-4-oxo-1,4-dihydro-3-
cinnolinecarboxamide;
N-(4-chlorobenzyl)-1-methyl-4-oxo-8-{4-[(4R)-2-oxo-1,3-
oxazolidin-4-yl]-1-butynyl}-6-(tetrahydro-2H-pyran-4-
ylmethyl) -1, 4-dihydro-3-cinnolinecarboxamide;
N-(4-chlorobenzyl)-8-(4-hydroxy-1-butynyl)-1-methyl-6-(4-hydroxy-1-butynyl)
morpholinylmethyl) -4-oxo-1, 4-dihydro-3-
cinnolinecarboxamide;
N-(4-chlorobenzyl)-8-[(1-hydroxycyclohexyl)ethynyl]-1-
methyl-6-(4-morpholinylmethyl)-4-oxo-1,4-dihydro-3-
cinnolinecarboxamide;
N-(4-chlorobenzyl)-8-(3,3-dicyclopropyl-3-hydroxy-1-
propynyl)-1-methyl-6-(4-morpholinylmethyl)-4-oxo-1,4-
dihydro-3-cinnolinecarboxamide;
N-(4-chlorobenzy1)-8-[(3S)-3-hydroxy-1-butyny1]-1-methyl-
6-(4-morpholinylmethyl)-4-oxo-1,4-dihydro-3-
cinnolinecarboxamide;
8-\{3-\{(aminocarbonyl)amino]-3-methyl-1-butynyl\}-N-(4-
chlorobenzyl)-1-methyl-6-(4-morpholinylmethyl)-4-oxo-1,4-
dihydro-3-cinnolinecarboxamide;
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N-(4-\text{chlorobenzyl})-1-\text{methyl}-8-[3-\text{methyl}-3-(4-\text{thioxo-})]
1,3,5-triazinan-1-y1)-1-butyny1]-6-(4-morpholinylmethyl)-
4-oxo-1, 4-dihydro-3-cinnolinecarboxamide;
N-(4-\text{chlorobenzyl})-8-[(3R)-3-\text{hydroxy}-1-\text{butynyl}]-1-\text{methyl}-
6-(4-morpholinylmethyl)-4-oxo-1,4-dihydro-3-
cinnolinecarboxamide:
N-(4-chlorobenzyl)-1-methyl-6-(4-morpholinylmethyl)-4-
oxo-8-{4-[(4R)-2-oxo-1,3-oxazolidin-4-yl]-1-butynyl}-1,4-
dihydro-3-cinnolinecarboxamide;
N-(4-\text{chlorobenzyl})-8-[3-(1,1-\text{dioxido}-4-\text{thiomorpholinyl})-
1-propynyl]-1-methyl-6-(4-morpholinylmethyl)-4-oxo-1,4-
dihydro-3-cinnolinecarboxamide;
N-(4-chlorobenzyl)-8-(5-hydroxy-1-pentynyl)-1-methyl-6-
(4-morpholinylmethyl)-4-oxo-1,4-dihydro-3-
cinnolinecarboxamide;
N-(4-chlorobenzyl)-8-\{[(1R,2S)-2-
hydroxycyclopentyl]ethynyl}-1-methyl-6-(4-
morpholinylmethyl) -4-oxo-1, 4-dihydro-3-
cinnolinecarboxamide;
N-(4-chlorobenzyl)-8-(3-hydroxy-3-methyl-1-butynyl)-1-
methyl-6-(4-morpholinylmethyl)-4-oxo-1,4-dihydro-3-
cinnolinecarboxamide:
N-(4-chlorobenzyl)-8-[3-(4,5-dichloro-1H-imidazol-1-yl)-
1-propynyl]-1-methyl-6-(4-morpholinylmethyl)-4-oxo-1,4-
dihydro-3-cinnolinecarboxamide;
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N-(4-chlorobenzyl)-8-(3-hydroxy-1-propynyl)-1-methyl-6-
(4-morpholinylmethyl)-4-oxo-1,4-dihydro-3-
cinnolinecarboxamide;
N-(4-chlorobenzyl)-1-methyl-4-oxo-8-(phenylethynyl)-6-
(tetrahydro-2H-pyran-4-ylmethyl)-1,4-dihydro-3-
cinnolinecarboxamide;
N-(4-chlorobenzyl)-8-(3-hydroxy-3-phenyl-1-propynyl)-1-
methyl-4-oxo-6-(tetrahydro-2H-pyran-4-ylmethyl)-1,4-
dihydro-3-cinnolinecarboxamide;
N-(4-chlorobenzyl)-8-(3-hydroxy-1-propynyl)-1-methyl-4-
oxo-1,4-dihydro-3-cinnolinecarboxamide;
N-(4-chlorobenzyl)-8-(4-hydroxy-1-butynyl)-1-methyl-4-
oxo-1,4-dihydro-3-cinnolinecarboxamide;
N-(4-chlorobenzyl)-8-(3-hydroxy-1-propynyl)-1-methyl-4-
oxo-6-(tetrahydro-2H-pyran-4-ylmethyl)-1,4-dihydro-3-
cinnolinecarboxamide;
N-(4-chlorobenzyl)-8-(4-hydroxy-1-butynyl)-1-methyl-4-
oxo-6-(tetrahydro-2H-pyran-4-ylmethyl)-1,4-dihydro-3-
cinnolinecarboxamide;
N-(4-chlorobenzyl)-8-[3-(dimethylamino)-1-propynyl]-1-
methyl-4-oxo-6-(tetrahydro-2H-pyran-4-ylmethyl)-1,4-
dihydro-3-cinnolinecarboxamide;
N-(4-chlorobenzyl)-1-[3-(methylsulfonyl)propyl]-6-(4-
morpholinylmethyl) -4-oxo-1, 4-dihydro-3-
cinnolinecarboxamide;
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N-(4-chlorobenzyl)-1-[3-(methylsulfanyl)propyl]-6-(4-
morpholinylmethyl)-4-oxo-1,4-dihydro-3-
cinnolinecarboxamide;
N-(4-\text{chlorobenzyl})-1-[(2-\text{hydroxyethoxy})\text{methyl}]-6-(4-\text{methyl})
morpholinylmethyl)-4-oxo-1,4-dihydro-3-
cinnolinecarboxamide;
N-(4-chlorobenzyl)-6-(4-morpholinylmethyl)-4-oxo-1-
tetrahydro-3-furanyl-1,4-dihydro-3-cinnolinecarboxamide;
N-(4-chlorobenzyl)-1-(1,2-diethyl-4-pyrazolidinyl)-6-(4-
morpholinylmethyl) -4-oxo-1, 4-dihydro-3-
cinnolinecarboxamide;
N-(4-chlorobenzyl)-6-(4-morpholinylmethyl)-1-(3-
oxetanyl) -4-oxo-1, 4-dihydro-3-cinnolinecarboxamide;
N-(4-chlorobenzyl)-1-{3-[(3-
hydroxypropyl)sulfonyl]propyl}-6-(4-morpholinylmethyl)-4-
oxo-1, 4-dihydro-3-cinnolinecarboxamide;
N-(4-chlorobenzyl)-1-[2-(2-ethoxyethoxy)ethyl]-6-(4-ethoxyethoxy)
morpholinylmethyl) -4-oxo-1, 4-dihydro-3-
cinnolinecarboxamide;
N-(4-chlorobenzyl)-6-(4-morpholinylmethyl)-4-oxo-1-
[(phenylsulfinyl)methyl]-1,4-dihydro-3-
cinnolinecarboxamide;
N-(4-\text{chlorobenzyl})-6-(4-\text{morpholinylmethyl})-4-\text{oxo}-1-
[(phenylsulfonyl)methyl]-1,4-dihydro-3-
cinnolinecarboxamide:
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N-(4-chlorobenzyl)-6-(4-morpholinylmethyl)-4-oxo-1-
[(phenylsulfanyl)methyl]-1,4-dihydro-3-
cinnolinecarboxamide;
N-(4-chlorobenzyl)-6-(4-morpholinylmethyl)-4-oxo-1-
tetrahydro-2H-pyran-3-yl-1,4-dihydro-3-
cinnolinecarboxamide:
N-(4-chlorobenzyl)-1-[(methylsulfanyl)methyl]-6-(4-
morpholinylmethyl)-4-oxo-1,4-dihydro-3-
cinnolinecarboxamide:
6-(4-morpholinylmethyl)-4-oxo-1,4-dihydro-3-
cinnolinecarboxamide;
N-(4-chlorobenzyl)-6-(4-morpholinylmethyl)-4-oxo-1-
tetrahydro-2H-pyran-4-yl-1,4-dihydro-3-
cinnolinecarboxamide;
N-(4-chlorobenzyl)-1-methyl-6-(4-morpholinylmethyl)-4-
oxo-8-(4-thiomorpholinylmethyl)-1,4-dihydro-3-
cinnolinecarboxamide:
N-(4-chlorobenzyl)-8-[(4-hydroxy-1-piperidinyl)methyl]-1-
methyl-6-(4-morpholinylmethyl)-4-oxo-1,4-dihydro-3-
cinnolinecarboxamide:
N-(4-chlorobenzyl)-8-\{[(3R)-3-
hydroxypyrrolidinyl]methyl}-1-methyl-6-(4-
morpholinylmethyl)-4-oxo-1,4-dihydro-3-
cinnolinecarboxamide:
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N-(4-chlorobenzyl)-8-[(3-hydroxy-1-piperidinyl)methyl]-1-
methyl-6-(4-morpholinylmethyl)-4-oxo-1,4-dihydro-3-
cinnolinecarboxamide;
[3-{[(4-chlorobenzyl)amino]carbonyl}-1-methyl-6-(4-
morpholinylmethyl)-4-oxo-1,4-dihydro-8-cinnolinyl]methyl
4-morpholinecarboxylate;
N-(4-chlorobenzyl)-8-(hydroxymethyl)-1-methyl-6-(4-
morpholinylmethyl) -4-oxo-1, 4-dihydro-3-
cinnolinecarboxamide:
N-(4-chlorobenzyl)-8-[(3-cyanobenzyl)amino]-1-methyl-6-
(4-morpholinylmethyl)-4-oxo-1,4-dihydro-3-
cinnolinecarboxamide;
N-(4-chlorobenzyl)-1-methyl-6,8-bis(4-morpholinylmethyl)-
4-oxo-1, 4-dihydro-3-cinnolinecarboxamide;
8-[(1-acetyl-4-piperidinyl)amino]-N-(4-chlorobenzyl)-1-
methyl-6-(4-morpholinylmethyl)-4-oxo-1,4-dihydro-3-
cinnolinecarboxamide;
N-(4-chlorobenzyl)-1-methyl-8-{[1-methyl-2-
(phenylsulfonyl)ethyl]amino}-6-(4-morpholinylmethyl)-4-
oxo-1,4-dihydro-3-cinnolinecarboxamide;
N-(4-chlorobenzyl)-8-\{[3-(4-methoxyphenyl)-1-
methylpropyl]amino}-1-methyl-6-(4-morpholinylmethyl)-4-
oxo-1, 4-dihydro-3-cinnolinecarboxamide;
8-amino-N-(4-chlorobenzyl)-1-methyl-6-(4-
morpholinylmethyl)-4-oxo-1,4-dihydro-3-
cinnolinecarboxamide;
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```
N-(4-chlorobenzyl)-1-methyl-6-(4-morpholinylmethyl)-8-
[(3-nitrobenzyl)amino]-4-oxo-1,4-dihydro-3-
cinnolinecarboxamide;
N-(4-chlorobenzyl)-1-methyl-6-(4-morpholinylmethyl)-4-
oxo-8-(tetrahydro-2H-pyran-4-ylamino)-1,4-dihydro-3-
cinnolinecarboxamide:
N-(4-chlorobenzyl)-6-(3-hydroxy-1-propyl)-1-methyl-4-oxo-
1,4-dihydro-3-cinnolinecarboxamide;
N-(4-chlorobenzyl)-6-(4-hydroxy-1-butyl)-1-methyl-4-oxo-
1,4-dihydro-3-cinnolinecarboxamide;
N-(4-\text{chlorobenzyl})-8-\{[(1R,2R)-1-\text{hydroxy}-2-
methylcyclohexyl]ethyl}-1-methyl-4-oxo-6-(tetrahydro-2H-
pyran-4-ylmethyl)-1,4-dihydro-3-cinnolinecarboxamide;
N-(4-chlorobenzyl)-8-(cyclopropylethyl)-1-methyl-6-(4-
morpholinylmethyl)-4-oxo-1,4-dihydro-3-
cinnolinecarboxamide:
N-(4-chlorobenzyl)-8-[3-(dimethylamino)-1-propyl]-1-
methyl-6-(4-morpholinylmethyl)-4-oxo-1,4-dihydro-3-
cinnolinecarboxamide;
N-(4-chlorobenzyl)-1-methyl-4-oxo-8-{4-[(4R)-2-oxo-1,3-
oxazolidin-4-yl]-1-butyl}-6-(tetrahydro-2H-pyran-4-
ylmethyl)-1,4-dihydro-3-cinnolinecarboxamide;
N-(4-\text{chlorobenzyl})-8-(4-\text{hydroxy-}1-\text{butyl})-1-\text{methyl-}6-(4-\text{methyl-}6)
morpholinylmethyl)-4-oxo-1,4-dihydro-3-
cinnolinecarboxamide;
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```
N-(4-chlorobenzyl)-8-[(1-hydroxycyclohexyl)ethyl]-1-
methyl-6-(4-morpholinylmethyl)-4-oxo-1,4-dihydro-3-
cinnolinecarboxamide;
N-(4-chlorobenzyl)-8-(3,3-dicyclopropyl-3-hydroxy-1-
propyl)-1-methyl-6-(4-morpholinylmethyl)-4-oxo-1,4-
dihydro-3-cinnolinecarboxamide;
N-(4-chlorobenzyl)-8-[(3S)-3-hydroxy-1-butyl]-1-methyl-6-
(4-morpholinylmethyl)-4-oxo-1,4-dihydro-3-
cinnolinecarboxamide;
8-{3-[(aminocarbonyl)amino]-3-methyl-1-butyl}-N-(4-
chlorobenzyl)-1-methyl-6-(4-morpholinylmethyl)-4-oxo-1,4-
dihydro-3-cinnolinecarboxamide;
N-(4-chlorobenzyl)-1-methyl-8-[3-methyl-3-(4-thioxo-
1,3,5-triazinan-1-yl)-1-butyl]-6-(4-morpholinylmethyl)-4-
oxo-1,4-dihydro-3-cinnolinecarboxamide;
N-(4-\text{chlorobenzyl})-8-[(3R)-3-\text{hydroxy}-1-\text{butyl}]-1-\text{methyl}-6-
(4-morpholinylmethyl)-4-oxo-1,4-dihydro-3-
cinnolinecarboxamide;
N-(4-chlorobenzyl)-1-methyl-6-(4-morpholinylmethyl)-4-
0x0-8-\{4-[(4R)-2-0x0-1,3-0xazolidin-4-yl]-1-butyl\}-1,4-
dihydro-3-cinnolinecarboxamide;
N-(4-chlorobenzyl)-8-[3-(1,1-dioxido-4-thiomorpholinyl)-
1-propyl]-1-methyl-6-(4-morpholinylmethyl)-4-oxo-1,4-
dihydro-3-cinnolinecarboxamide;
N-(4-chlorobenzyl)-8-(5-hydroxy-1-pentyl)-1-methyl-6-(4-
morpholinylmethyl)-4-oxo-1,4-dihydro-3-
cinnolinecarboxamide;
```

```
N-(4-chlorobenzyl)-8-\{[(1R,2S)-2-
hydroxycyclopentyl]ethyl}-1-methyl-6-(4-
morpholinylmethyl) -4-oxo-1, 4-dihydro-3-
cinnolinecarboxamide;
N-(4-chlorobenzyl)-8-(3-hydroxy-3-methyl-1-butyl)-1-
methyl-6-(4-morpholinylmethyl)-4-oxo-1,4-dihydro-3-
cinnolinecarboxamide;
N-(4-\text{chlorobenzyl})-8-[3-(4,5-\text{dichloro-1H-imidazol-1-yl})-
1-\text{propyl}] -1-\text{methyl} -6-(4-\text{morpholinylmethyl}) -4-\text{oxo} -1, 4-\text{oxo}
dihydro-3-cinnolinecarboxamide;
N-(4-chlorobenzyl)-8-[3-(1H-imidazol-1-yl)-1-propyl]-1-
methyl-6-(4-morpholinylmethyl)-4-oxo-1,4-dihydro-3-
cinnolinecarboxamide;
N-(4-\text{chlorobenzyl})-8-[3-(1H-\text{imidazol}-1-\text{yl})-1-\text{propynyl}]-1-
methyl-6-(4-morpholinylmethyl)-4-oxo-1,4-dihydro-3-
cinnolinecarboxamide;
N-(4-chlorobenzyl)-8-(3-hydroxy-1-propyl)-1-methyl-6-(4-
morpholinylmethyl) -4-oxo-1, 4-dihydro-3-
cinnolinecarboxamide:
N-(4-chlorobenzyl)-1-methyl-4-oxo-8-(phenylethyl)-6-
(tetrahydro-2H-pyran-4-ylmethyl)-1,4-dihydro-3-
cinnolinecarboxamide;
N-(4-chlorobenzyl)-8-(3-hydroxy-3-phenyl-1-propyl)-1-
methyl-4-oxo-6-(tetrahydro-2H-pyran-4-ylmethyl)-1,4-
dihydro-3-cinnolinecarboxamide;
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N-(4-chlorobenzyl)-8-(3-hydroxy-1-propyl)-1-methyl-4-oxo-1,4-dihydro-3-cinnolinecarboxamide;

N-(4-chlorobenzyl)-8-(4-hydroxy-1-butyl)-1-methyl-4-oxo-1,4-dihydro-3-cinnolinecarboxamide;

N-(4-chlorobenzyl)-8-(3-hydroxy-1-propyl)-1-methyl-4-oxo-6-(tetrahydro-2H-pyran-4-ylmethyl)-1,4-dihydro-3-cinnolinecarboxamide;

N-(4-chlorobenzyl)-8-(4-hydroxy-1-butyl)-1-methyl-4-oxo-6-(tetrahydro-2H-pyran-4-ylmethyl)-1,4-dihydro-3-cinnolinecarboxamide;

N-(4-chlorobenzyl)-8-[3-(dimethylamino)-1-propyl]-1-methyl-4-oxo-6-(tetrahydro-2H-pyran-4-ylmethyl)-1,4-dihydro-3-cinnolinecarboxamide;

N-(4-chlorobenzyl)-1-methyl-8-{[methyl(tetrahydro-2-furanylmethyl)amino]methyl}-6-(4-morpholinylmethyl)-4-oxo-1,4-dihydro-3-cinnolinecarboxamide;

and pharmaceutically acceptable salts thereof.

5. The method of Claim 1, wherein the compound administered has the Formula VII

VII

or a pharmaceutically acceptable salt thereof, wherein,

 $A^{VII}$  is

- a) Cl,
- b) Br,
- c) CN,
- d)  $NO_2$ , or
- e) F;

#### $R^{VII-1}$ is

- a) aryl<sup>VII</sup>,
- b)  $S(0)_{m}^{VII}R^{VII-6}$ ,
- c) (C=0) $R^{VII-6}$ , with the proviso that if  $R^{VII-6}$  is  $NR^{VII-7}R^{VII-8}$ , then  $R^{VII-7}$  and  $R^{VII-8}$  do not both equal H
- d) (C=O)  $OR^{VII-9}$ ,
- e) cyano,
- f)  $\text{het}^{\text{VII}}$ , wherein said  $\text{het}^{\text{VII}}$  is bound via a carbon atom,
- g) Ohet<sup>VII</sup>,
- h)  $NR^{VII-7}R^{VII-8}$  with the proviso that  $R^{VII-7}$  and  $R^{VII-8}$  do not both equal H
- i)  $SR^{VII-10}$ ,
- j) Shet<sup>VII</sup>,
- k) NHCOR<sup>VII-12</sup>,
- 1)  $NHSO_2R^{VII-12}$ ,
- m)  $C_{1-7}$ alkyl which is partially unsaturated and optionally substituted by one or more substituents of the group  $R^{VII-11}$ ,  $OR^{VII-13}$ ,  $SR^{VII-10}$ ,  $SR^{VII-13}$ ,  $NR^{VII-7}R^{VII-8}$ , halo,  $(C=0)C_{1-7}$ alkyl, or  $SO_mR^{VII-9}$ , or
- n)  $C_{1-7}$ alkyl which is substituted by one or more substituents of the group  $R^{\text{VII-11}}$ ,  $OR^{\text{VII-13}}$ ,  $SR^{\text{VII-10}}$ ,  $SR^{\text{VII-13}}$ ,  $NR^{\text{VII-7}}R^{\text{VII-8}}$ , halo,  $(C=O)C_{1-7}$ alkyl, or  $SO_m^{\text{VII}}R^{\text{VII-9}}$ ;

# $R^{VII-2}$ is

- a) H,
- b) halo,
- c) aryl<sup>VII</sup>,

- d)  $S(0)_{m}^{VII}R^{VII-6}$ ,
- e) (C=O)  $R^{VII-6}$ ,
- f) (C=O)  $OR^{VII-9}$ ,
- g) cyano,
- h)  $\text{het}^{\text{VII}}$ , wherein said  $\text{het}^{\text{VII}}$  is bound via a carbon atom,
- i) OR $^{VII-10}$ ,
- j) Ohet<sup>VII</sup>,
- k)  $NR^{VII-7}R^{VII-8}$ .
- 1)  $SR^{VII-10}$ ,
- m) Shet<sup>VII</sup>,
- n) NHCOR<sup>VII-12</sup>,
- o)  $NHSO_2R^{VII-12}$ , or
- p)  $C_{1-7}$ alkyl which may be partially unsaturated and optionally substituted by one or more substituents of the group  $R^{VII-11}$ ,  $OR^{VII-13}$ ,  $SR^{VII-10}$ ,  $SR^{VII-13}$ ,  $NR^{VII-7}R^{VII-8}$ , halo,  $(C=0)C_{1-7}$ alkyl, or  $SO_m^{VII}R^{VII-9}$ , or
- q)  $R^{\text{VII-1}}$  together with  $R^{\text{VII-2}}$  form a carbocyclic or het which may be optionally substituted by  $NR^{\text{VII-7}}R^{\text{VII-8}}$ , or  $C_{1-7}$  alkyl which may be optionally substituted by  $OR^{\text{VII-14}}$ ;

# $R^{VII-6}$ is

- a)  $C_{1-7}$ alkyl,
- b)  $NR^{VII-7}R^{VII-8}$
- c) aryl<sup>VII</sup>, or
- d) het<sup>VII</sup>, wherein said het<sup>VII</sup> is bound via a carbon atom;

# $\textbf{R}^{\text{VII--7}}$ and $\textbf{R}^{\text{VII--8}}$ are independently

- a) H,
- b) aryl<sup>VII</sup>,
- c)  $C_{1-7}$ alkyl which may be partially unsaturated and is optionally substituted by one or more substituents selected from  $NR^{VII-10}R^{VII-10}$ ,  $R^{VII-11}$ ,  $SO_mR^{VII-9}$ ,  $CONR^{VII-10}R^{VII-10}$ , or halo, or,

d)  $R^{VII-7}$  and  $R^{VII-8}$  together with the nitrogen to which they are attached form a het  $^{VII}$ ;

# $R^{VII-9}$ is

- a) aryl<sup>VII</sup>,
- b) het<sup>VII</sup>,
- c)  $C_{3-8}$ cycloalkyl,
- d) methyl, or
- e)  $C_{2-7}$ alkyl which may be partially unsaturated and is optionally substituted by one or more substituents selected from  $NR^{VII-10}R^{VII-10}$ ,  $R^{VII-11}$ , SH,  $CONR^{VII-10}R^{VII-10}$ , or halo;

## $R^{VII-10}$ is

- a) H,
- b) methyl, or
- c) C<sub>2-7</sub>alkyl optionally substituted by OH;

## R<sup>VII-11</sup> is

- a)  $OR^{VII-10}$ ,
- b) Ohet VII,
- c) Oaryl<sup>VII</sup>,
- d)  $CO_2R^{VII-10}$ ,
- e) het<sup>VII</sup>,
- f) aryl<sup>VII</sup>,
- g) CN, or
- h)  $C_{3-8}$ cycloalkyl which may be partially unsaturated and optionally substituted by one or more substituents seleted from a group consisting of  $R^{\text{VII-11}}$ ,  $NR^{\text{VII-7}}R^{\text{VII-8}}$ ,  $SO_m^{\text{VII}}R^{\text{VII-9}}$ , or  $C_{1-7}$ alkyl optionally substituted by  $R^{\text{VII-11}}$ ,  $NR^{\text{VII-7}}R^{\text{VII-8}}$ , or  $SO_m^{\text{VII}}R^{\text{VII-9}}$ ;

# $R^{VII-12}$ is

- a) H,
- b) het<sup>VII</sup>,
- c) aryl<sup>VII</sup>,
- d)  $C_{3-8}$ cycloalkyl,
- e) methyl, or

f)  $C_{2-7}$ alkyl optionally substituted by  $NR^{VII-7}R^{VII-8}$  or  $R^{VII-11}$ ;

#### R<sup>VII-13</sup> is

- a)  $(P=0) (OR^{VII-14})_{2}$
- b)  $CO(CH_2)_n^{VII}CON(CH_3) (CH_2)_nSO_3^{-M^{VII+}}$ ,
- c) an amino acid,
- d)  $C(=0) \operatorname{aryl}^{VII}$ , or
- e)  $C(=O)C_{1-7}alkyl$  optionally substituted by  $NR^{VII-7}R^{VII-8}$ ,  $aryl^{VII}$ ,  $het^{VII}$ ,  $CO_2H$ , or  $O(CH_2)_n^{VII}CO_2R^{VII-14}$ ;

## $R^{VII-14}$ is

- a) H, or
- b)  $C_{1-7}$ alkyl;

each  $n^{VII}$  is independently 1, 2, 3, 4 or 5; each  $m^{VII}$  is independently 0, 1, or 2;  $M^{VII}$  is sodium, potassium, or lithium;

- aryl is a phenyl radical or an ortho-fused bicyclic carbocyclic radical wherein at least one ring is aromatic;
- wherein any aryl<sup>VII</sup> is optionally substituted with one or more substituents selected from the group consisting of halo, OH, cyano,  $CO_2R^{VII-14}$ ,  $CF_3$ ,  $C_{1-6}$ alkoxy, and  $C_{1-6}$  alkyl which may be further substituted by one to three  $SR^{VII-14}$ ,  $NR^{VII-14}R^{VII-14}$ ,  $OR^{VII-14}$ , or  $CO_2R^{VII-14}$  groups;
- het<sup>VII</sup> is a four- (4), five- (5), six- (6), or seven- (7) membered saturated or unsaturated heterocyclic ring having 1, 2, or 3 heteroatoms selected from the group consisting of oxygen, sulfur, and nitrogen, which is optionally fused to a benzene ring, or any bicyclic heterocycle group;
- wherein any het<sup>VII</sup> is optionally substituted with one or more substituents selected from the group consisting of halo, OH, cyano, phenyl,  $CO_2R^{VII-14}$ ,  $CF_3$ ,  $C_{1-6}$ alkoxy, oxo, oxime, and  $C_{1-6}$  alkyl which may be further

substituted by one to three  $SR^{VII-14}$ ,  $NR^{VII-14}R^{VII-14}$ ,  $OR^{VII-14}$ , or  $CO_2R^{VII-14}$  groups.

- 6. The method of Claim 5, wherein  $A^{VII}$  is Cl.
- 7. The method of Claim 6, wherein  $R^{VII-1}$  is selected from the group consisting of  $CH_2$ -morpholine, alkynl- $CH_2OH$ ,  $CH_2$ -(tetrahydro-2H-pyran-4-yl) and  $(CH_2)_3OH$ .
- 8. The compound of Claim 6, wherein the compound administered is selected from the group consisting of

N-(4-chlorobenzyl)-4-hydroxy-6-(tetrahydro-2H-pyran-4-ylmethyl)-3-cinnolinecarboxamide;

N-(4-chlorobenzyl)-4-hydroxy-6-(4-morpholinylmethyl)-3-cinnolinecarboxamide;

Methyl 3-{[(4-chlorobenzyl)amino]carbonyl}-4-hydroxy-6-cinnolinecarboxylate;

N-(4-chlorobenzyl)-4-hydroxy-6-(hydroxymethyl)-3-cinnolinecarboxamide <math>N-(4-chlorobenzyl)-8-(cyclopropylethynyl)-4-hydroxy-6-

(4-morpholinylmethyl)-3-cinnolinecarboxamide;

N-(4-chlorobenzyl)-8-[3-(dimethylamino)-1-propynyl]-4-hydroxy-6-(4-morpholinylmethyl)-3-cinnolinecarboxamide;

N-(4-chlorobenzyl)-4-hydroxy-8-(4-hydroxy-1-butynyl)-6-(4-morpholinylmethyl)-3-cinnolinecarboxamide;

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N-(4-chlorobenzyl)-4-hydroxy-8-[(1-
hydroxycyclohexyl)ethynyl]-6-(4-morpholinylmethyl)-3-
cinnolinecarboxamide;
N-(4-chlorobenzyl)-8-(3,3-dicyclopropyl-3-hydroxy-1-
propynyl) -4-hydroxy-6-(4-morpholinylmethyl) -3-
cinnolinecarboxamide;
N-(4-chlorobenzyl)-4-hydroxy-8-[(3S)-3-hydroxy-1-
butynyl]-6-(4-morpholinylmethyl)-3-cinnolinecarboxamide;
8-{3-[(aminocarbonyl)amino]-3-methyl-1-butynyl}-N-(4-
chlorobenzyl) -4-hydroxy-6-(4-morpholinylmethyl) -3-
cinnolinecarboxamide:
N-(4-chlorobenzyl)-4-hydroxy-8-[3-methyl-3-(4-thioxo-
1,3,5-triazinan-1-yl)-1-butynyl]-6-(4-morpholinylmethyl)
-3-cinnolinecarboxamide;
N-(4-chlorobenzyl)-4-hydroxy-8-[(3R)-3-hydroxy-1-
butynyl]-6-(4-morpholinylmethyl)-3-cinnolinecarboxamide;
N-(4-chlorobenzyl)-4-hydroxy-6-(4-morpholinylmethyl)-8-
\{4-[(4R)-2-oxo-1,3-oxazolidin-4-yl]-1-butynyl\}-3-
cinnolinecarboxamide;
N-(4-chlorobenzyl)-8-[3-(1,1-dioxido-4-thiomorpholinyl)-
1-propynyl]-4-hydroxy-6-(4-morpholinylmethyl)-3-
cinnolinecarboxamide;
N-(4-chlorobenzyl)-4-hydroxy-8-(5-hydroxy-1-pentynyl)-6-
(4-morpholinylmethyl)-3-cinnolinecarboxamide;
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N-(4-chlorobenzyl)-4-hydroxy-8-{(1R,2S)-2-}
hydroxycyclopentyl]ethynyl}-6-(4-morpholinylmethyl)-3-
cinnolinecarboxamide;
N-(4-chlorobenzyl)-4-hydroxy-8-(3-hydroxy-3-methyl-1-
butynyl)-6-(4-morpholinylmethyl)-3-cinnolinecarboxamide;
N-(4-\text{chlorobenzyl})-8-[3-(4,5-\text{dichloro-1H-imidazol-1-yl})-
1-propynyl]-4-hydroxy-6-(4-morpholinylmethyl)-3-
cinnolinecarboxamide:
N-(4-chlorobenzyl)-4-hydroxy-8-(3-hydroxy-1-propynyl)-6-
(4-morpholinylmethyl)-3-cinnolinecarboxamide;
N-(4-chlorobenzyl)-8-(cyclopropylethyl)-4-hydroxy-6-(4-
morpholinylmethyl) -3-cinnolinecarboxamide;
N-(4-chlorobenzyl)-8-[3-(dimethylamino)-1-propyl]-4-
hydroxy-6-(4-morpholinylmethyl)-3-cinnolinecarboxamide;
N-(4-\text{chlorobenzyl})-4-\text{hydroxy}-8-(4-\text{hydroxy}-1-\text{butyl})-6-(4-\text{hydroxy}-1-\text{hydroxy})
morpholinylmethyl) -3-cinnolinecarboxamide;
N-(4-chlorobenzyl)-4-hydroxy-8-[(1-
hydroxycyclohexyl)ethyl]-6-(4-morpholinylmethyl)-3-
cinnolinecarboxamide;
N-(4-chlorobenzyl)-8-(3,3-dicyclopropyl-3-hydroxy-1-
propyl)-4-hydroxy-6-(4-morpholinylmethyl)-3-
cinnolinecarboxamide;
N-(4-chlorobenzy1)-4-hydroxy-8-[(3S)-3-hydroxy-1-buty1]-
6-(4-morpholinylmethyl)-3-cinnolinecarboxamide;
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chlorobenzyl)-4-hydroxy-6-(4-morpholinylmethyl)-3-
cinnolinecarboxamide;
N-(4-\text{chlorobenzyl})-4-\text{hydroxy-8-}[3-\text{methyl-3-}(4-\text{thioxo-})]
1,3,5-triazinan-1-y1)-1-butyl]-6-(4-morpholinylmethyl)-3-
cinnolinecarboxamide:
N-(4-chlorobenzyl)-4-hydroxy-8-[(3R)-3-hydroxy-1-butyl]-
6-(4-morpholinylmethyl)-3-cinnolinecarboxamide;
N-(4-chlorobenzyl)-4-hydroxy-6-(4-morpholinylmethyl)-8-
\{4-[(4R)-2-oxo-1,3-oxazolidin-4-yl]-1-butyl\}-3-
cinnolinecarboxamide:
N-(4-chlorobenzyl)-8-[3-(1,1-dioxido-4-thiomorpholinyl)-
1-propyl]-4-hydroxy-6-(4-morpholinylmethyl)-3-
cinnolinecarboxamide:
N-(4-chlorobenzyl)-4-hydroxy-8-(5-hydroxy-1-pentyl)-6-(4-
morpholinylmethyl) -3-cinnolinecarboxamide;
N-(4-chlorobenzyl)-4-hydroxy-8-{[(1R,2S)-2-
hydroxycyclopentyl]ethyl}-6-(4-morpholinylmethyl)-3-
cinnolinecarboxamide;
N-(4-\text{chlorobenzyl})-4-\text{hydroxy-8}-(3-\text{hydroxy-3-methyl-1}-
butyl)-6-(4-morpholinylmethyl)-3-cinnolinecarboxamide;
N-(4-\text{chlorobenzyl})-8-[3-(4,5-\text{dichloro-1H-imidazol-1-yl})-
1-propyl]-4-hydroxy-6-(4-morpholinylmethyl)-3-
cinnolinecarboxamide;
N-(4-\text{chlorobenzyl})-4-\text{hydroxy-8}-(3-\text{hydroxy-1-propyl})-6-(4-\text{chlorobenzyl})
morpholinylmethyl)-3-cinnolinecarboxamide;
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 $8-\{3-[(aminocarbonyl)amino]-3-methyl-1-butyl\}-N-(4-$ 

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N-(4-chlorobenzyl)-4-hydroxy-8-(3-hydroxy-1-propynyl)-3-
cinnolinecarboxamide;
N-(4-chlorobenzyl)-4-hydroxy-8-(4-hydroxy-1-butynyl)-3-
cinnolinecarboxamide;
N-(4-chlorobenzyl)-4-hydroxy-8-(3-hydroxy-1-propynyl)-6-
(tetrahydro-2H-pyran-4-ylmethyl)-3-cinnolinecarboxamide;
N-(4-chlorobenzyl)-4-hydroxy-8-(phenylethynyl)-6-
(tetrahydro-2H-pyran-4-ylmethyl)-3-cinnolinecarboxamide;
N-(4-chlorobenzyl)-4-hydroxy-8-(3-hydroxy-3-phenyl-1-
propynyl)-6-(tetrahydro-2H-pyran-4-ylmethyl)-3-
cinnolinecarboxamide:
N-(4-chlorobenzyl)-4-hydroxy-8-(4-hydroxy-1-butynyl)-6-
(tetrahydro-2H-pyran-4-ylmethyl)-3-cinnolinecarboxamide;
N-(4-chlorobenzyl)-8-[3-(dimethylamino)-1-propynyl]-4-
hydroxy-6-(tetrahydro-2H-pyran-4-ylmethyl)-3-
cinnolinecarboxamide:
N-(4-chlorobenzyl)-4-hydroxy-8-{[(1R,2R)-1-hydroxy-2-
methylcyclohexyl]ethynyl}-6-(tetrahydro-2H-pyran-4-
ylmethyl)-3-cinnolinecarboxamide;
N-(4-chlorobenzyl)-4-hydroxy-8-{4-(4R)-2-oxo-1,3-}
oxazolidin-4-yl]-1-butynyl}-6-(tetrahydro-2H-pyran-4-
ylmethyl) -3-cinnolinecarboxamide;
N-(4-chlorobenzyl)-4-hydroxy-8-(3-hydroxy-1-propyl)-6-
(tetrahydro-2H-pyran-4-ylmethyl)-3-cinnolinecarboxamide;
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N-(4-chlorobenzyl)-4-hydroxy-8-(phenylethyl)-6-(tetrahydro-2H-pyran-4-ylmethyl)-3-cinnolinecarboxamide;

N-(4-chlorobenzyl)-4-hydroxy-8-(3-hydroxy-3-phenyl-1-propyl)-6-(tetrahydro-2H-pyran-4-ylmethyl)-3-cinnolinecarboxamide;

N-(4-chlorobenzyl)-4-hydroxy-8-(4-hydroxy-1-butyl)-6-(tetrahydro-2H-pyran-4-ylmethyl)-3-cinnolinecarboxamide;

N-(4-chlorobenzyl)-8-[3-(dimethylamino)-1-propyl]-4-hydroxy-6-(tetrahydro-2H-pyran-4-ylmethyl)-3-cinnolinecarboxamide;

N-(4-chlorobenzyl)-4-hydroxy-8-{[(1R,2R)-1-hydroxy-2-methylcyclohexyl]ethyl}-6-(tetrahydro-2H-pyran-4-ylmethyl)-3-cinnolinecarboxamide;

N-(4-chlorobenzyl)-4-hydroxy-8-{4-[(4R)-2-oxo-1,3-oxazolidin-4-yl]-1-butyl}-6-(tetrahydro-2H-pyran-4-ylmethyl)-3-cinnolinecarboxamide;

and pharmaceutically acceptable salts thereof.

9. A method of Claim 1, wherein the compound administered is Formula VIII

and pharmaceutically acceptable salts thereof, wherein  $\mathbf{A}^{\text{VIII}}$  is

- Cl, a) b) Br, CN, C) d)  $NO_2$ , or F; e)  $R^{VIII-1}$  is R<sup>VIII-5</sup>, a) NR<sup>VIII-7</sup>R<sup>VIII-8</sup>, or b) SO<sub>2</sub>R<sup>VIII-9</sup>; C)  $R^{VIII-2}$  is aryl<sup>VIII</sup>, a) het<sup>VIII</sup>, b) SOm VIIIR VIII-6, C) d)
  - $OC_{2-7}$  alkyl substituted by OH,
  - $SC_{2-7}$  alkyl substituted by OH, or e)
  - f)  $C_{2-8}$  alkyl which is partially unsaturated and is optionally substituted by one or more substituents selected from R<sup>VIII-11</sup>, OR<sup>VIII-13</sup>,  $\text{SR}^{\text{VIII-13}},~\text{NR}^{\text{VIII-7}}\text{R}^{\text{VIII-8}},~\text{halo,}~\text{(C=O)C}_{\text{1-7}}~\text{alkyl}~\text{or}$ SOm VIII R VIII-9;

with the proviso that when  $R^{\text{VIII-1}} = R^{\text{VIII-5}} =$ (CH2CH2O)  $_{i}R^{\text{VIII-10}}$ , then  $R^{\text{VIII-2}}$  may additionally represent

- Η, a)
- b) halo,
- (C=O) R<sup>VIII-6</sup>, C)
- (C=O) OR<sup>VIII-9</sup>, d)
- e) cyano,
- OR<sup>VIII-10</sup>, f)
- Ohet VIII, g)
- NR<sup>VIII-7</sup>R<sup>VIII-8</sup>, h)
- SR<sup>VIII-10</sup>, i)
- Shet<sup>VIII</sup>, j)
- NHCOR VIII-12, k)

- 1)  $NHSO_2R^{VIII-12}$ , or
- m)  $R^{VIII-2}$  together with  $R^{VIII-3}$  or  $R^{VIII-4}$  form a carbocyclic or het which may be optionally substituted by  $NR^{VIII-7}R^{VIII-8}$ , or  $C_{1-7}$ alkyl which may be optionally substituted by  $OR^{VIII-14}$ ;

 $R^{VIII-3}$  and  $R^{VIII-4}$  are independently:

- a) H,
- b) halo,
- c) aryl<sup>VIII</sup>,
- d)  $S(0)_{m}^{VIII}R^{VIII-6}$ ,
- e)  $(C=0) R^{VIII-6}$ ,
- f) (C=O)  $OR^{VIII-9}$ ,
- g) cyano,
- h) het<sup>VIII</sup>, wherein said het<sup>VIII</sup> is bound via a carbon atom,
- i) OR<sup>VIII-10</sup>,
- j) Ohet<sup>VIII</sup>,
- k)  $R^{VIII-7}R^{VIII-8}$
- 1) SR<sup>VIII-10</sup>,
- m) Shet<sup>VIII</sup>,
- n) NHCOR VIII-12,
- o) NHSO<sub>2</sub>R<sup>VIII-12</sup>,
- p)  $C_{1-7}$ alkyl which may be partially unsaturated and optionally substituted by one or more substituents of the group  $R^{VIII-11}$ ,  $OR^{VIII-13}$ ,  $SR^{VIII-10}$ ,  $SR^{VIII-13}$ ,  $NR^{VIII-7}R^{VIII-8}$ , halo,  $(C=0)C_{1-7}$  alkyl, or  $SO_m^{VIII}R^{VIII-9}$ , or
- q)  $R^{VIII-4}$  together with  $R^{VIII-3}$  form a carbocyclic or het which may be optionally substituted by  $NR^{VIII-7}R^{VIII-8}$ , or  $C_{1-7}$  alkyl which may be optionally substituted by  $OR^{VIII-14}$ ;

 $R^{VIII-5}$  is

- a)  $(CH_2CH_2O)_i^{VIII}R^{VIII-10}$ ,
- b) het<sup>VIII</sup>, wherein said het<sup>VIII</sup> is bound via a carbon atom,
- c) aryl<sup>VIII</sup>,
- d)  $C_{1-7}$ alkyl which may be partially unsaturated and is optionally substituted by one or more substituents selected from  $NR^{VIII-7}R^{VIII-8}$ ,  $R^{VIII-11}$ ,  $SO_m^{VIII}R^{VIII-9}$ , or  $OC_{2-4}$ alkyl which may be further substituted by het  $^{VIII}$ ,  $OR^{VIII-10}$ , or  $NR^{VIII-7}R^{VIII-8}$ , or
- e)  $C_{3-8}$ cycloalkyl which may be partially unsaturated and optionally substituted by one or more substituents selected from  $R^{VIII-11}$ ,  $NR^{VIII-7}R^{VIII-8}$ ,  $SO_m^{VIII}R^{VIII-9}$ , or  $C_{1-7}$ alkyl optionally substituted by  $R^{VIII-11}$ ,  $NR^{VIII-7}R^{VIII-8}$ , or  $SO_mR^{VIII-9}$ ;

## $R^{VIII-6}$ is

- a)  $C_{1-7}$ alkyl,
- b) NR<sup>VIII-7</sup>R<sup>VIII-8</sup>,
- c) aryl<sup>VIII</sup>, or
- d) het<sup>VIII</sup>, wherein said het<sup>VIII</sup> is bound via a carbon atom;

# $R^{VIII-7}$ and $R^{VIII-8}$ are independently

- a) H,
- b) aryl<sup>VIII</sup>,
- C)  $C_{1-7}$ alkyl which may be partially unsaturated and is optionally substituted by one or more substituents selected from  $NR^{VIII-10}R^{VIII-10}$ ,  $R^{VIII-11}$ ,  $SO_mR^{VIII-9}$ ,  $CONR^{VIII-10}R^{VIII-10}$ , or halo, or,
- d)  $R^{VIII-7}$  and  $R^{VIII-8}$  together with the nitrogen to which they are attached form a het  $^{VIII}$ ;

## R<sup>VIII-9</sup> is

a) aryl<sup>VIII</sup>,

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het<sup>VIII</sup>,
       b)
       C)
              C_{3-8}cycloalkyl,
              methyl, or
       d)
              C_{2-7}alkyl which may be partially unsaturated and
       e)
               is optionally substituted by one or more
               substituents selected from NR^{VIII-10}R^{VIII-10},
               R^{VIII-11}, SH, CONR^{VIII-10}R^{VIII-10}, or halo;
R<sup>VIII-10</sup> is
               Η,
       a)
               methyl, or
       b)
               C_{2-7}alkyl optionally substituted by OH;
       C)
R^{VIII-11} is
               OR<sup>VIII-10</sup>,
       a)
               Ohet^{\text{VIII}},
       b)
               Oaryl<sup>VIII</sup>,
       C)
               CO_2R^{VIII-10},
       d)
              het<sup>VIII</sup>,
        e)
               aryl^{VIII}, or
        f)
        g)
               CN;
R^{VIII-12} is
        a)
               Η,
              het<sup>VIII</sup>,
        b)
               aryl<sup>VIII</sup>,
        C)
              C_{3-8}cycloalkyl,
        d)
              methyl, or
        e)
               \text{C}_{\text{2-7}}alkyl optionally substituted by \text{NR}^{\text{VIII-7}}R^{\text{VIII-8}}
        f)
               or R<sup>VIII-11</sup>;
R^{VIII-13} is
                (P=0) (OR^{14})_2,
        a)
               CO(CH_2)_n^{VIII}CON(CH_3) - (CH_2)_nSO_3^-M^{VIII+},
        b)
              an amino acid,
        C)
              C(=0)aryl<sup>VIII</sup>, or
        d)
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e)  $C(=O)C_{1-7}alkyl$  optionally substituted by  $NR^{VIII-7}R^{VIII-8}$ ,  $aryl^{VIII}$ ,  $het^{VIII}$ ,  $CO_2H$ , or  $O(CH_2)_n^{VIII}CO_2R^{VIII-14}$ ;

#### R<sup>VIII-14</sup> is

- a) H, or
- b)  $C_{1-7}$ alkyl;

each i<sup>VIII</sup> is independently 2, 3, or 4; each n<sup>VIII</sup> is independently 1, 2, 3, 4 or 5; each m<sup>VIII</sup> is independently 0, 1, or 2; M<sup>VIII</sup> is sodium, potassium, or lithium; aryl<sup>VIII</sup> is a phenyl radical or an ortho-fused bicyclic carbocyclic radical wherein at least one ring is aromatic;

- wherein any aryl<sup>VIII</sup> is optionally substituted with one or more substituents selected from halo, OH, cyano,  $CO_2R^{VIII-14}$ ,  $CF_3$ ,  $C_{1-6}$ alkoxy, and  $C_{1-6}$  alkyl which may be further substituted by one to three  $SR^{VIII-14}$ ,  $NR^{VIII-14}R^{VIII-14}$ ,  $OR^{VIII-14}$ , or  $CO_2R^{VIII-14}$  groups;
- het<sup>VIII</sup> is a four- (4), five- (5), six- (6), or seven- (7) membered saturated or unsaturated heterocyclic ring having 1, 2, or 3 heteroatoms selected from the group consisting of oxygen, sulfur, and nitrogen, which is optionally fused to a benzene ring, or any bicyclic heterocycle group;
- wherein any het<sup>VIII</sup> is optionally substituted with one or more substituents selected from the group consisting of halo, OH, cyano, phenyl,  $CO_2R^{VIII-14}$ ,  $CF_3$ ,  $C_{1-6}$ alkoxy, oxo, oxime, and  $C_{1-6}$  alkyl which may be further substituted by one to three  $SR^{VIII-14}$ ,  $NR^{VIII-14}R^{VIII-14}$ ,  $OR^{VIII-14}$ , or  $CO_2R^{VIII-14}$  groups.

- 10. The method of Claim 9, wherein  $A^{VIII}$  is Cl.
- 11. The method of Claim 9, wherein  $R^{\text{VIII-2}}$  is alkynl-CH<sub>2</sub>OH.
- 12. The method of Claim 9, wherein the compound administered is N-(4-chlorobenzyl)-6-(3-hydroxy-1-propynyl)-1,7-dimethyl-4-oxo-1,4-dihydro[1,8]naphthyridine-3-carboxamide, or N-(4-chlorobenzyl)-6-(3-hydroxy-1-propynyl)-7-methoxy-1-methyl-4-oxo-1,4-dihydro[1,8]naphthyridine-3-carboxamide; or a pharmaceutically acceptable salt thereof.
- 13. The method of Claim 9, wherein the compound administered is:

N-(4-chlorobenzyl)-6-(3-hydroxy-1-propynyl)-1,7-dimethyl-4-oxo-1,4-dihydro[1,8]naphthyridine-3-carboxamide;

N-(4-chlorobenzyl)-6-(3-hydroxypropyl)-1,7-dimethyl-4-oxo-1,4-dihydro[1,8]naphthyridine-3-carboxamide;

N-(4-Chlorobenzyl)-6-iodo-7-methoxy-1-methyl-4-oxo-1,4-dihydro[1,8]naphthyridine-3-carboxamide;

N-(4-chlorobenzyl)-1,7-dimethyl-6-(4-morpholinylmethyl)-4-oxo-1,4-dihydro[1,8]naphthyridine-3-carboxamide;

N-(4-chlorobenzyl)-1-methyl-4,7-dioxo-1,4,7,8-tetrahydro[1,8]naphthyridine-3-carboxamide;

N-(4-chlorobenzyl)-6-(3-hydroxy-1-propynyl)-7-methoxy-1- methyl-4-oxo-1,4-dihydro[1,8]naphthyridine-3-carboxamide;

N-(4-chlorobenzyl)-6-(3-hydroxypropyl)-7-methoxy-1-methyl-4-oxo-1,4-dihydro[1,8]naphthyridine-3-carboxamide;

ethyl 6-{[(4-chlorobenzyl)amino]carbonyl}-2-methoxy-8-methyl-5-oxo-5,8-dihydro[1,8]naphthyridine-3-carboxylate;

and pharmaceutically acceptable salts thereof.

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and pharmaceutically acceptable salts thereof, wherein,  $\mathbf{R}^{\mathrm{IX-1}}$  is

- a) Cl,
- b) Br,
- c) CN,
- d)  $NO_2$ , or
- e) F;

 $\textbf{R}^{\text{IX-2}},~\textbf{R}^{\text{IX-3}}$  and  $\textbf{R}^{\text{IX-4}}$  are independently selected from:

- a) H,
- b) halo,
- c) aryl<sup>IX</sup>,
- d)  $S(0)_{m}^{IX}R^{IX-6}$ ,
- e)  $(C=0) R^{IX-6}$ ,
- f) (C=O)  $OR^{IX-9}$ ,

- g) cyano,
- h)  $\text{het}^{\text{IX}}$ , wherein said  $\text{het}^{\text{IX}}$  is bound via a carbon atom,
- i) OR<sup>IX-10</sup>,
- j) Ohet<sup>IX</sup>,
- k)  $NR^{IX-7}R^{IX-8}$
- 1)  $SR^{IX-10}$ ,
- m) S<sup>IX-</sup>het,
- n) NHCOR<sup>IX-12</sup>,
- o)  $NHSO_2R^{IX-12}$ , or
- p)  $C_{1-7}$ alkyl which may be partially unsaturated and optionally substituted by one or more substituents of the group  $R^{IX-11}$ ,  $OR^{IX-13}$ ,  $SR^{IX-10}$ ,  $SR^{IX-13}$ ,  $NR^{IX-7}R^{IX-8}$ , halo,  $(C=0)C_{1-7}$ alkyl, or  $SO_mR^{IX-9}$ ;

## $R^{IX-6}$ is

- a)  $C_{1-7}$ alkyl,
- b)  $NR^{IX-7}R^{IX-8}$ ,
- c) aryl<sup>IX</sup>, or
- d)  $het^{IX}$ , wherein said  $het^{IX}$  is bound via a carbon atom;

# $R^{\text{IX-7}}$ and $R^{\text{IX-8}}$ are independently

- a) H,
- b) aryl<sup>IX</sup>,
- C)  $C_{1-7}$ alkyl which may be partially unsaturated and is optionally substituted by one or more substituents selected from  $NR^{IX-10}R^{IX-10}$ ,  $R^{IX-11}$ ,  $SO_mR^{IX-9}$ ,  $CONR^{IX-10}R^{IX-10}$ , or halo, or,
- d)  $R^{IX-7}$  and  $R^{IX-8}$  together with the nitrogen to which they are attached form a het<sup>IX</sup>;

## R<sup>IX-9</sup> is

a) aryl<sup>IX</sup>,

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C)
               C_{3-8}cycloalkyl,
       d)
               methyl, or
               C_{2-7}alkyl which may be partially unsaturated and
       e)
               is optionally substituted by one or more
               substituents selected from NR^{IX-10}R^{IX-10}, R^{IX-11},
               SH, CONR<sup>IX-10</sup>R<sup>IX-10</sup>, or halo;
R^{IX-10} is
               Η,
       a)
       b)
               methyl, or
               C_{2-7}alkyl optionally substituted by OH;
R^{\text{IX-11}} is
               ORIX-10,
       a)
       b)
               Ohet<sup>IX</sup>,
               Oaryl<sup>IX</sup>,
       C)
               CO_2R^{IX-10},
       d)
              het<sup>IX</sup>,
       e)
               aryl<sup>IX</sup>, or
       f)
               CN;
       g)
R^{\text{IX-12}} is
       a)
               Η,
               het<sup>IX</sup>,
       b)
               aryl<sup>IX</sup>,
       C)
       d)
               C_{3-8}cycloalkyl,
               methyl, or
       e)
               C_{2\text{--7}}alkyl optionally substituted by NR^{\text{IX--7}}R^{\text{IX--8}} or
       f)
               R<sup>IX-11</sup>;
R^{\text{IX-13}} is
               (P=0) (OR^{IX-14})_{2}
       a)
               CO(CH_2)_n^{IX}CON(CH_3) - (CH_2)_n^{IX}SO_3^{-M}^{IX+},
       b)
       C)
               an amino acid,
       d)
             C(=0) aryl, or
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het<sup>IX</sup>,

b)

- e)  $C(=0)C_{1-7}alkyl$  optionally substituted by  $NR^{IX-7}R^{IX-8}, \ aryl^{IX}, \ het^{IX}, \ CO_2H, \ or \ O(CH_2)_nCO_2R^{IX-14};$   $R^{IX-14}$  is
  - a) H, or
  - b)  $C_{1-7}$ alkyl;

each n<sup>IX</sup> is independently 1, 2, 3, 4 or 5; each m<sup>IX</sup> is independently 0, 1, or 2; M<sup>IX</sup> is sodium, potassium, or lithium;

- aryl is a phenyl radical or an ortho-fused bicyclic carbocyclic radical wherein at least one ring is aromatic;
- wherein any aryl<sup>IX</sup> is optionally substituted with one or more substituents selected from the group consisting of halo, OH, cyano,  $CO_2R^{IX-14}$ ,  $CF_3$ ,  $C_{1-6}$ alkoxy, and  $C_{1-6}$  alkyl which may be further substituted by one to three  $SR^{IX-14}$ ,  $NR^{IX-14}R^{IX-14}$ ,  $OR^{IX-14}$ , or  $CO_2R^{IX-14}$  groups;
- het<sup>IX</sup> is a four- (4), five- (5), six- (6), or seven- (7) membered saturated or unsaturated heterocyclic ring having 1, 2, or 3 heteroatoms selected from the group consisting of oxygen, sulfur, and nitrogen, which is optionally fused to a benzene ring, or any bicyclic heterocycle group;
- wherein any het<sup>IX</sup> is optionally substituted with one or more substituents selected from the group consisting of halo, OH, cyano, phenyl,  $CO_2R^{IX-14}$ ,  $CF_3$ ,  $C_{1-6}$ alkoxy, oxo, oxime, and  $C_{1-6}$  alkyl which may be further substituted by one to three  $SR^{IX-14}$ ,  $NR^{IX-14}R^{IX-14}$ ,  $OR^{IX-14}$ , or  $CO_2R^{IX-14}$  groups.

- 15. The method of Claim 14, wherein  $R^{IX-1}$  is Cl.
- 16. The method of Claim 14, wherein the compound administered is selected from a group consisting of

N-(4-chlorobenzyl)-4-hydroxy-7-methyl[1,8]naphthyridine-3-carboxamide;

N-(4-chlorobenzyl)-4-hydroxy-7-methyl-6-(tetrahydro-2H-pyran-4-ylmethyl)[1,8]naphthyridine-3-carboxamide;

N-(4-chlorobenzyl)-4-hydroxy-7-methyl-6-(4-morpholinylmethyl)[1,8]naphthyridine-3-carboxamide;

6-bromo-N-(4-chlorobenzyl)-4-hydroxy-7-methyl[1,8]naphthyridine-3-carboxamide;

N-(4-chlorobenzyl)-4-hydroxy-6-(3-hydroxy-1-propynyl)-7-methyl[1,8]naphthyridine-3-carboxamide;

N-(4-chlorobenzyl)-4-hydroxy-6-iodo-7-methyl[1,8]naphthyridine-3-carboxamide; and

Methyl 6-{[(4-chlorobenzyl)amino]carbonyl}-5-hydroxy-2-methyl[1,8]naphthyridine-3-carboxylate.

- 17. The method according to Claim 1, wherein said mammal is a human.
- 18. The method according to Claim 1, wherein said mammal is a livestock or companion animal.
- 19. The method according to Claim 1, wherein the amount administered is from about 0.1 to about 300 mg/kg of mammal body weight.

- 20. The method according to Claim 1, wherein the amount administered is from about 1 to about 30 mg/kg of mammal body weight.
- 21. The method according to Claim 2, wherein the compound is administered parenterally, intravaginally, intranasally, topically, orally, or rectally.